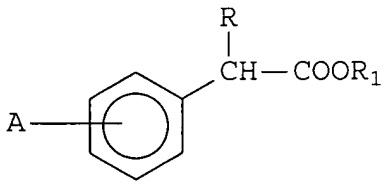
AMENDMENTS TO THE CLAIMS

This listing of claims replaces all prior versions, and listings, of claims in the present application.

IN THE CLAIMS:

1. (Currently Amended) A process for the preparation of meta or para-substituted α-arylalkanoic acids of formula (I):



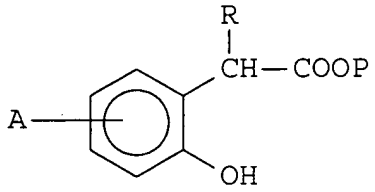
(I)

wherein[:]

R is hydrogen; C_1-C_6 alkyl; R_1 is hydrogen, straight or branched C_1-C_6 alkyl, phenyl, p-nitrophenyl, a cation of an alkali or alkaline-earth metal cation or of a pharmaceutically acceptable ammonium salt; A is C_1-C_4 alkyl, aryl, aryloxy, arylcarbonyl, 2-, 3- or 4-pyridocarbonyl, aryl optionally substituted with one or more alkyl, hydroxy, amino, cyano, nitro, alkoxy, haloalkyl, haloalkoxy; A is at the meta or para positions;

which process comprises the following steps:

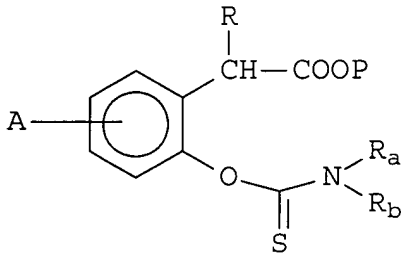
a) ~~transformation~~ reaction of compounds of formula (II)



(II)

in which P is straight or branched C_1-C_6 alkyl, phenyl, p-nitrophenyl,

~~into~~ with a thiocarbonyl halide to give compounds of formula (III)

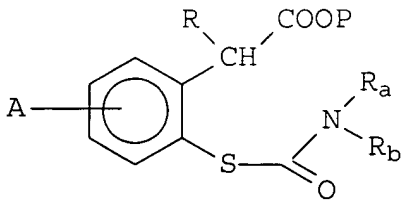


(III)

wherein

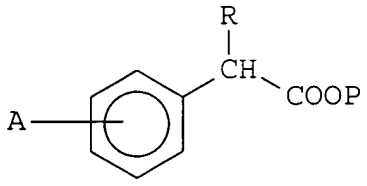
R_a and R_b are C_1-C_6 alkyl ,

b) thermal rearrangement of compound (III) to give (IIIb)



(IIIb)

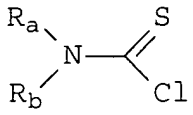
c) catalytic hydrogenation of (IIIb) to give (IIIc)



(IIIc)

d) ~~transformation~~ hydrolysis of (IIIc) and optional subsequent reesterfication or salification to give ~~into~~ (I).

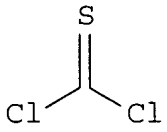
2. (Original) A process according to claim 1, in which the transformation of step a) is carried out by reaction of the compound (II) with



wherein R_a and R_b are as defined in claim 1, in the presence of an organic or inorganic base.

3. (Original) A process as claimed in claim 2, in which said organic base is selected from triethylamine and pyridine, and said inorganic base is selected from alkali or alkaline-earth carbonates.

4. (Currently Amended) A process as claimed in claim 1, in which the transformation of step a) is carried out by reaction of compound (II) with thiophosgene

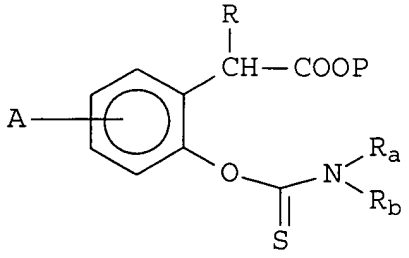


and the subsequent reaction of the resulting product with HNR_aR_b, wherein R_a and R_b are as defined in claim 1.

5. (Original) A process as claimed in claim 1, in which the hydrogenation of step c) is carried out with Ni-Raney.

6. (Original) A process according to any one of the above claims, in which the group A of formula (I) is meta-benzoyl and R is methyl.

7. (Currently Amended) As a reaction intermediate, the compound

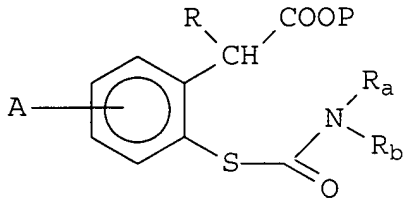


(III)

wherein [:]

R is hydrogen, C_1-C_6 alkyl; A is a C_1-C_4 alkyl, aryl, aryloxy, aryl optionally substituted with one or more alkyl, hydroxy, amino, cyano, nitro, alkoxy, haloalkyl, haloalkoxy, A is at the meta or para positions; P is straight or branched C_1-C_6 alkyl, phenyl, p-nitrophenyl; R_a and R_b are C_1-C_6 alkyl.

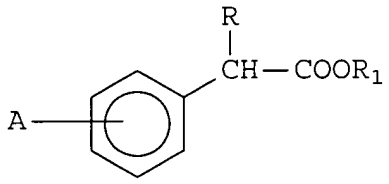
8. (Original) As a reaction intermediate, the compound



(IIIb)

wherein A, R, P, R_a and R_b are as defined in claim 7.

9. (New) A process for the preparation of meta or para-substituted α-arylalkanoic acids of formula (I):

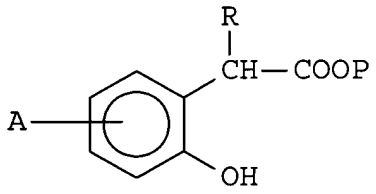


(I)

wherein R is hydrogen; C_1-C_6 alkyl; R_1 is hydrogen, straight or branched C_1-C_6 alkyl, phenyl, p-nitrophenyl, a cation of an alkali or alkaline-earth metal cation or of a pharmaceutically acceptable ammonium salt; A is C_1-C_4 alkyl, aryl, aryloxy, arylcarbonyl, 2-, 3- or 4- pyridocarbonyl, aryl optionally substituted with one or more alkyl, hydroxy, amino, cyano, nitro, alkoxy, haloalkyl, haloalkoxy; A is at the meta or para positions;

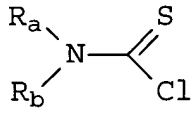
wherein said process comprises the steps of:

a) reaction of compounds of formula (II)



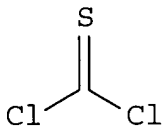
(II)

in which P is straight or branched C_1-C_6 alkyl, phenyl, p-nitrophenyl, with the compound

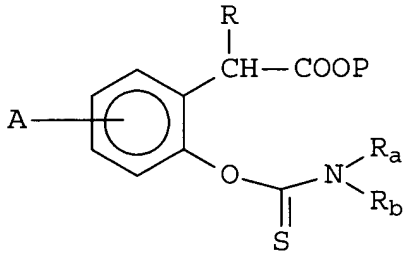


wherein R_a and R_b are C_1-C_6 alkyl, in the presence of an organic or inorganic base, or

with a thiophosgene



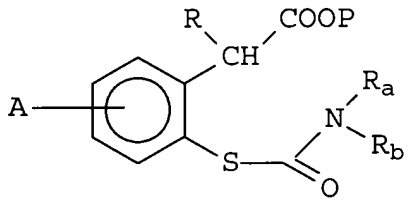
and the subsequent reaction of the resulting product with HNR_aR_b, wherein R_a and R_b are as defined above, to give compounds of formula (III)



(III)

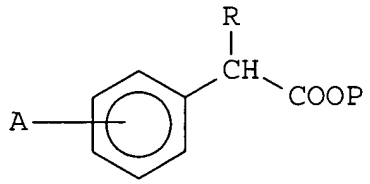
wherein R_a and R_b are as defined above,

b) thermal rearrangement of compound (III) to give (IIIb)



(IIIb)

c) catalytic hydrogenation of (IIIb) to give (IIIc)



(IIIc)

d) hydrolysis of (IIIc) and optional subsequent reesterfication or salification to give (I).